

Video Article

An Experimental Model to Study Tuberculosis-Malaria Coinfection upon Natural Transmission of *Mycobacterium tuberculosis* and *Plasmodium berghei*

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URL: <http://www.jove.com/video/50829>

DOI: [doi:10.3791/50829](https://doi.org/10.3791/50829)

Keywords: Infectious Diseases, Issue 84, coinfection, mouse, Tuberculosis, Malaria, *Plasmodium berghei*, *Mycobacterium tuberculosis*, natural transmission

Date Published: 2/17/2014

Citation: Mueller, A.K., Behrends, J., Blank, J., Schaible, U.E., Schneider, B.E. An Experimental Model to Study Tuberculosis-Malaria Coinfection upon Natural Transmission of *Mycobacterium tuberculosis* and *Plasmodium berghei*. *J. Vis. Exp.* (84), e50829, doi:10.3791/50829 (2014).

Abstract

Coinfections naturally occur due to the geographic overlap of distinct types of pathogenic organisms. Concurrent infections most likely modulate the respective immune response to each single pathogen and may thereby affect pathogenesis and disease outcome. Coinfected patients may also respond differentially to anti-infective interventions. Coinfection between tuberculosis as caused by mycobacteria and the malaria parasite *Plasmodium*, both of which are coendemic in many parts of sub-Saharan Africa, has not been studied in detail. In order to approach the challenging but scientifically and clinically highly relevant question how malaria-tuberculosis coinfection modulate host immunity and the course of each disease, we established an experimental mouse model that allows us to dissect the elicited immune responses to both pathogens in the coinfecting host. Of note, in order to most precisely mimic naturally acquired human infections, we perform experimental infections of mice with both pathogens by their natural routes of infection, i.e. aerosol and mosquito bite, respectively.

Video Link

The video component of this article can be found at <http://www.jove.com/video/50829/>

Introduction

Human populations are rarely exposed to one pathogen only. Particularly in regions with high incidence of infections such as sub-Saharan Africa, coinfections represent a prime but highly underappreciated public health problem. Tuberculosis and malaria are the most prevalent bacterial and parasitic infections in humans, respectively and continue to be major causes of morbidity and mortality in impoverished populations in the tropics. Despite the wide geographic overlap between tuberculosis and malaria and the large number of individuals at risk of coinfection, very little is known about the interactions between the various and often counteracting immune regulators and effectors simultaneously elicited against malaria parasites and tubercle bacilli in coinfecting individuals.

Rodent models of mixed infections allow characterizing differential immune reactions against distinct pathogens in one host, thus shedding light on the mutual influences modulating pathology and clinical outcome of the individual disease, which will also help to identify new recommendations for therapy and prevention. We have established an experimental mouse model which enables us to investigate the ramifications caused by concurrent infection with *Mycobacterium tuberculosis* (*Mtb*) and rodent *Plasmodium* species within the same host¹. Importantly, to emulate naturally acquired human infections as closely as possible, our model implements the natural routes of infection used by both pathogens. Tuberculosis is an airborne infection, which therefore primarily manifests in the lungs. The causative agent, *Mtb* is transmitted by people with active disease via the aerosol route when they expel infectious aerosol droplets during coughing or sneezing. Thus, aerosol infection is the method of choice to mimic natural infection. Generation of airborne particles that contain *Mtb* can be achieved by the use of an inhalation exposure system. This whole body exposure chamber allows exposing experimental animals to *Mtb*-containing infectious aerosol droplets (**Figure 1**) which are inhaled into the alveoli of the lungs where infection is initiated².

Malaria on the other hand is a vector-borne disease caused by the protozoan, apicomplexan parasite *Plasmodium* that is naturally transmitted by the bite of a female Anopheline mosquito. During the blood meal, infectious sporozoites are deposited under the host skin and subsequently reach the liver via the blood stream. Within hepatocytes they develop and rapidly multiply into liver-stage schizonts. Eventually, mature schizonts rupture and release thousands of pathogenic first-generation merozoites into the blood stream where they initiate a progressive cycle of red cell invasion, replication, red cell rupture, and reinvasion³. The *Plasmodium* life cycle continues as some merozoites develop into the sexual parasite stages, the male and female gametocytes, which can be taken up by mosquitoes during blood meals. In the mosquito, *Plasmodium* sporozoites develop within oocysts residing in the mosquito midgut and eventually migrate to the salivary glands for onward transmission into another host^{3,4}.

While in experimental animal models the delivery of *Mtb* via the aerosol and thus most relevant route is very common, many experimental rodent *Plasmodium* infection studies including studies on malaria-tuberculosis coinfection⁵⁻⁷ have been carried out by infecting mice with parasitized erythrocytes, giving rise to blood-stage malaria infection while excluding the clinically-silent liver-stage phase. However, the liver stage is an obligatory step during infection and relevant for anti-plasmodial immunity⁸⁻¹¹. We consider it therefore important to include the hepatic phase of malaria transmission and maintenance in studies of both malaria and malaria-tuberculosis coinfection, respectively. In addition, it has been shown that naturally transmitted sporozoites are more infectious than those transmitted via needle infections¹², which prompted us to establish the malaria infection by mosquito bite instead of injecting isolated salivary gland derived sporozoites. The only way to obtain infectious mosquitoes for natural transmission of malaria parasites is to maintain the whole life cycle of the parasite in both the vertebrate host (here mouse) and the mosquito vector. Thus, access to an insectary for parasite maintenance is inevitable in order to perform natural transmission *by bite*.

Our protocol described herein was developed to investigate how coinfection with *Plasmodium* sporozoites impacts on chronic tuberculosis¹. To do so, mice are aerosol infected with *M. tuberculosis* and 40 days later, when *M. tuberculosis* infection has reached the chronic phase, mice are exposed to malaria-infectious mosquitoes. The outcome of both malaria and tuberculosis in the coinfecting animals can be followed by monitoring blood parasitemia and bacterial load in tissue, respectively. In our protocol, we will describe in detail how to infect mice with both pathogens via their natural infection route and how to confirm successful pathogen transmission. In our hands, the commonly achieved infection rate for both experimental infections is 100%. These protocols can be applied to study both infections separately or, as we do, to model and study coinfection between two of the most challenging human infectious diseases. This model is applicable to other *Mycobacterium* and rodent *Plasmodium* species beyond the ones described in this protocol.

Protocol

Safety Measures and Ethics Statement

The studies presented herein involve work with the human pathogen *Mtb* and the rodent malaria parasite *Plasmodium berghei* (*P. berghei*). Experiments with *Mtb* (strain H37Rv) and *P. berghei* must be carried out under the appropriate biosafety conditions. Biosafety level (BSL) 2 laboratories are required for rodent malaria parasites such as *P. berghei* and BSL 3 for *Mtb* and hence, for all coinfection studies described herein. NOTE: Mice infected with *Mtb* do not spread the infection to other mice within immediate vicinity (our own observation). Spread to the experimenter can therefore be excluded; however appropriate protective clothing has to be worn inside the BSL 3 laboratory at all times. According to our rules, experimenters wear surgical scrubs, hair nets, disposable respirators, and two pairs of gloves, of which the outer one is discarded anytime arms are withdrawn from the cabinet. New ones are put on before entering the cabinet again. Two containers, one with 2% Buraton (NOTE: other disinfectants approved to inactivate *Mtb* can be used at the approved concentration) for liquid and infectious waste material and one for surface decontamination, are prepared before the experiment and placed inside the cabinet. In addition, plastic bags will be used for solid noninfectious waste. According to German rules, all work involving the handling and processing of *Mtb* cultures or tissue derived from *Mtb* infected mice must be carried out in a class 2 biosafety cabinet within a BSL3 lab. While handling mycobacteria, measures have to be taken to avoid the generation of aerosols at all times.

Female C57BL/6 mice (Charles River) aged between 6-8 weeks were used for all coinfection experiments and maintained under specific barrier conditions in BSL 3 facilities. Animal care and experimentation were performed in accordance with protocols approved by the Ethics Committee for Animal Experiments of the Ministry for Agriculture, Environment, and Rural Areas of the State of Schleswig-Holstein

For obtaining malarial mosquito stages, NMRI mice were purchased from Charles River Laboratory, Sulzfeld, Germany and kept under specific pathogen-free conditions within the University Heidelberg animal facility (IBF). All animal experiments were performed according to European regulations and approved by the state authorities of Baden-Württemberg (Regierungspräsidium Karlsruhe).

1. Aerosol Infection of Mice with *Mtb* using a Glas-Col Aerosol Chamber

The best way to standardize aerosol infection of experimental animals with *Mtb* is to use mycobacteria from frozen stocks with known CFU titers. To prepare stock cultures, culture *Mtb* in Middlebrook 7H9 broth supplemented with OADC (Oleic acid, Albumin, Dextrose, Catalase) enrichment medium and 0.05% Tween 80, a carbon source for mycobacteria and measure to avoid bacterial clumping at the same time. Cultures should have an OD ≤ 1 at the time of harvest. At higher OD, bacterial clumping increases and viability decreases. Store 1 ml aliquots at -80 °C. Determine the number of viable colony forming units (CFU) in frozen stocks by plating a series of 10-fold dilutions of three independent vials on 7H11 agar plates supplemented with 0.5% glycerol, 1 g/L asparagine, and OADC and enumerate colonies after 4 weeks of incubation at 37 °C. Perform aerosol infection as described below.

1. Thaw *Mtb* stocks of known CFU titer and carefully mix the suspension five times to disperse bacterial clumps using a 1 ml syringe fitted with a 27 G needle. Avoid the production of aerosols.
2. Depending on the desired infection dose, transfer the required volume of the mycobacterial stock into a 50 ml tube containing sterile PBS. The final volume is 6 ml. NOTE: By varying the number of microorganism in this suspension, the proportion of bacteria bearing aerosol droplets is varied. We usually aim at an uptake of 100 viable bacilli per lung (low-dose infection). In our experience, this requires around 1-2 x 10⁶ *Mtb*/ml in a total volume of 6 ml of which 5.5 ml are nebulized (see below). It is recommended to do a series of experimental aerosol challenges with different concentrations of bacteria to find the ideal conditions for your infection. Whereas high dose infections with up to 5,000 mycobacteria can be done to speed up the infectious process, as few as 5 bacteria can be used to successfully infect mice by aerosol. The infection rate is commonly 100%.
3. Remove sufficient volume (prepared in excess of final volume required) for plating to determine the inoculum titer; we usually remove 500 µl to plate technical triplicates.

4. Place animals in a compartmented mesh basket (one mouse per basket) within the circular aerosol chamber and close the lid of the aerosol chamber. NOTE: Other models are equipped with a pie-shaped basket composed of five individual compartments, each of which can accommodate 20 mice.
5. Attach the Venturi-nebulizer unit to the three stainless steel socket joints.
6. Remove the mycobacterial suspension from the 50 ml tube with a 10 ml syringe fitted with an 18 G blunt needle and carry the syringe to the aerosol chamber in a closed transport box.
7. Remove the screw cap of the nebulizer unit and carefully inject the mycobacteria suspension into the nebulizer. Avoid the generation of aerosols. Discard the syringe into a sharps container containing 2% Buraton. Seal the nebulizer with the screw cap.
8. Switch on the main power switch and the UV lamp. The display on the control keypad shows "Glas-Col Apparatus Co".
9. Turn the program switch on. The display will show "Is the nebulizer ready?" "Is the basket loaded?" Press enter when ready". Press enter.
10. The display shows "Enter Preheat Time 900"; this means the preheat time for the incinerator (which decontaminates the exhaust air) is 900 sec. Press enter.
11. The display will show "Enter Nebulizing time 1,800"; this means the nebulizing time is set to 1,800 sec as default. In order to extend the nebulizing time, enter "2,400" and press enter. NOTE: During the nebulizing cycle, air under pressure atomizes the suspension, thereby generating small aerosol droplets containing mycobacteria. With the main air flow, these droplets (approximately 2-5 μm in size) are carried into the aerosol chamber.
12. The display will show "Enter C.D. Time 1,800"; this means that the decay cycle takes 1,800 sec. Set to "2,400" and press enter. NOTE: During this cycle, the cloud which has been built up in the aerosol chamber during the nebulizing cycle can decay. The small droplets are inhaled by the experimental animals.
13. The display will show "Enter Dec Time 900"; this means that the UV light decontamination cycle will take 900 sec. Press enter. The machine will start cycling through preheat, nebulizing, cloud decay and UV decontamination.

CAUTION: The vacuum flow meter should indicate 60 cubic feet/hr (check when preheat cycle starts; adjust vacuum control valve if necessary) and the compressed air flow meter 10 cubic feet/hr (check when nebulizing cycle starts; adjust air control valve accordingly).

14. When the cycle is complete, the keypad will show "Process Complete – Remove Specimen". Turn off the program, UV and main power switch.
15. Check if the mycobacterial suspension has been nebulized completely and if not, record the remaining volume by carefully removing the suspension with an appropriate syringe fitted with an 18 G blunt needle. The remaining volume should not be more than 1 ml.
16. Remove the nebulizer from the joints and place it in a pan containing 2% Buraton for a minimum of 2 hr, usually disinfection is done O/N. Afterwards, transfer the nebulizer to a fresh pan and rinse thoroughly with water. Leave nebulizer to air-dry.
17. Open the aerosol chamber and return animals to their cages. Bag the baskets in autoclave bags and autoclave. Wipe clean the surfaces of the inside of the aerosol chamber with 2% Buraton. NOTE: For safety reasons, a powered air purifying respirator should be worn during this procedure.
18. Using the remaining 500 μl of the mycobacterial suspension (see step 1.3) plate 10-fold dilutions of a technical triplicate (3 x 100 μl) on 7H11 agar plates.

2. Verify Uptake of Desired CFU in Lungs

One day after aerosol infection of experimental animals determine the bacterial load in the lungs of designated control animals in order to verify the uptake of the desired CFU. NOTE: We usually designate an extra group of 3-5 mice for day 1 CFU determination.

To monitor the course of *Mtb* infection over time, the mycobacterial tissue burden can be examined by plating serial dilutions of whole organ homogenates for CFU determination. The lung is the primary site for disease manifestation in tuberculosis however spleen, liver and lymph nodes are usually analyzed accordingly. The dilutions to be plated depend on the expected mycobacterial load in the organs, which depend on the initial inoculum (low dose vs. high dose), which gives rise to different bacterial loads in tissue, as well as on the organ and time point to be analyzed. Upon low dose infection, the bacterial load of *Mtb* H37Rv in the lung usually reaches a plateau between day 25-30.

1. Place euthanized animals (Euthanasia: CO₂ asphyxiation or terminal anesthesia) on absorbent paper on a dissecting board in a class II biosafety cabinet and disinfect mice with 70% ethanol. NOTE: 70% alcohol is for surface decontamination only and will not kill *Mtb*.
2. Make a small incision in the middle of the abdomen and retract the mouse skin above the head.
3. Open the abdomen and the thoracic cage with surgical scissors and remove the thoracic wall so that the lungs are accessible.
4. Remove the lungs and transfer into a 15 ml tube with homogenization buffer (sterile water/1% v/v Tween 80/1% w/v albumin).
5. Using the plunger of a 5 ml syringe strain lungs through 100 μm sieves into a small Petri dish.
6. Flush sieves several times using a 1 ml pipette equipped with a barrier tip and distribute the lung homogenate among 8 agar plates (ca. 250 μl /plate; make sure they have a well dried surface).
7. Carefully plate the samples using disposable spreaders which are discarded into 2% Buraton.
8. Leave agar plates to dry inside the cabinet. Seal every single plate with parafilm, wrap in aluminum foil and incubate upright at 37 °C for at least 4 weeks.

3. Construction of Escape-proof Mosquito Cages

Rodent *Plasmodium* strains are not harmful to humans. However, since *Mtb*-infected mice are the recipients of the parasite and work is done under BSL 3 conditions, precautions are necessary to prevent the mosquitoes from escaping their cages. Depending on the planned infection regimen, autoclavable and reusable metal-frame cages or self-made disposable cages can be used. The latter ones are recommended if *by bite* infection of experimental animals is required to be carried out by a defined number of mosquitoes per mouse. If self-made cages are required, prepare cages to exact specifications as the health of the mosquitoes and the safety of the laboratory depends upon their sound construction (Figure 2).

1. Take a carton such as a half-pint ice cream carton or paper coffee cup (Figure 2).

2. Cut a small opening into the side of the carton and cover with a double layer of latex with a slit cut in each piece to create a secure entrance for the mosquitoes. NOTE: The opening has an overall size of 2 cm x 2 cm, the tubing (aspirator device) that we use to bring in and/or take-out mosquitoes is a modified 15 ml Falcon tube attached to a vacuum pump and hence serves as an aspirator with a diameter of 1.5 cm.
3. Tape a piece of filter paper on the inside bottom of the carton to soak up drips.
4. Close off ice cream cups with netting (double layer of Nylon mesh, size 1 mm x 1 mm) and fix to the carton by tape and elastic band.
5. For infection experiments where defined numbers of mosquitoes per mouse are required, prepare cages with 10-15 infected mosquitoes, each of which ideally contains an average of 10,000 wild-type salivary gland *P. berghei* sporozoites.
6. Ideally, starve mosquitoes for 24 hr prior to performing the blood meal.

4. Malaria Infection of Mice by Mosquito Bite

The rodent malaria parasite used herein is *P. berghei* however any other rodent *Plasmodium* strain of interest can be used. To obtain infectious mosquitoes for sporozoite transmission the whole life cycle of the parasite is maintained in both the vertebrate host (mouse) and the mosquito vector. Parasite maintenance is carried out in an insectary. For natural transmission experiments, the minimum number of sporozoites per salivary gland should be no less than 10,000. For detailed protocols on parasite maintenance we refer to *Methods in Malaria Research* by Moll et al., 2013.

1. Anaesthetize naïve mice or animals preinfected for 40 days with *Mtb* with Ketamine (100 mg/kg) and Xylazine (7 mg/kg) solution by intraperitoneal injection (200 µl/mouse). NOTE: The time between *Mtb* and *Plasmodium* infection can be adjusted depending on the underlying research question. Likewise, the order of pathogen challenges can be reversed, i.e. mice can be exposed to infectious mosquito bite prior to *Mtb* infection.
2. Place anaesthetized mice onto the netting of the mosquito cages (one mouse per mosquito cage for defined mosquito to mouse ratio) and allow the mosquitoes to feed through the membrane for 10-15 min. NOTE: The presence of blood in the mosquito guts implies feeding and thus, sporozoite transmission.
3. Transfer mice back into their cages and supervise constantly until they wake up from anesthesia. NOTE: Place mice on paper towel and not on bedding (risk of suffocation) and keep warm (place closely together and cover bodies with paper towel).
4. Kill the mosquitoes by spraying them with 70% ethanol or other disinfectants through the netting of the cage. Autoclave cages and discard if disposable ones were used.

5. Monitoring Parasitemia

1. Puncture tail veins at the very end with a needle and collect one drop of blood at each end of a slide.
2. Use another slide to pull one drop of blood over half of the slide's length. Flip over the spreader to use the other edge for the second smear. Leave slides to air dry.
3. Giemsa stain
 1. Place slides in slide holders and fix the blood smears by brief immersion in absolute methanol. NOTE: Because the use of glass ware should be kept at minimum in the BSL 3 we use polypropylene cuvettes for all solutions.
 2. Immerse the smears in Giemsa (1:10 in deionized water). After 10 min, dip the slides in and out of staining cuvettes filled with deionized water a few times to rinse away the excess stain.
 3. Finally, air dry slides in a vertical position.
4. ALTERNATIVE STAIN: Wright's stain
 1. Dissolve Wright's stain at a concentration of 1 mg/ml in methanol.
 2. Filter through folded filter prior to use.
 3. Immerse the blood smears in staining solution and incubate for 4 min (NOTE: methanol fixation of smears is not necessary as the staining solution is methanol based).
 4. Wash by rinsing slides in deionized water as described above and leave slides to air dry.
 5. Once Giemsa/Wright's stain is complete, gently transfer all reagents used into a Buraton-containing disposal pot.
5. Blood smear analysis
 1. Analyze stained blood smears by light microscopy at a 100-fold magnification with oil immersion.
 2. Count uninfected and infected erythrocytes in an area of the blood smear where erythrocytes are arranged in a monolayer.
 3. In order to achieve a well-defined parasitemia, count at least 10 different fields of view displaying an erythrocyte monolayer.
 4. Calculate the relative parasitemia (in %) by dividing the number of parasitized erythrocytes by the number of erythrocytes and multiplying by 100.

Representative Results

Infection of mice with *Mtb* and *P. berghei* via the natural route, i.e. aerosol and mosquito bite, respectively, results in consistent and reproducible infection of all animals (Table 1)¹. We can monitor successful infection and the course of both malaria and tuberculosis by determining the initial occurrence (prepatency) and number of parasites in the blood (*P. berghei* parasitemia) and the burden of *Mtb* in different organs, respectively. An example of successful malaria parasite transmission by infectious mosquito bite is shown in Table 1 and Figure 3. Feeding of 10 sporozoite-infected mosquitoes on each mouse resulted in a 100% infection rate as confirmed by the presence of blood-stage parasites 4-5 days later (Table 1 and Figure 3A). Importantly, we find similar parasite numbers in the blood of individual mice of the same experimental group, indicating that sporozoite transmission by mosquito bite results in a consistent and reproducible infection. By comparing the prepatency in naïve control mice with that in *Mtb*-coinfected animals, we can see that prepatency is slightly delayed in mice that had been preinfected with *Mtb* (Table 1, Figure 3B)¹. As

we have reported previously¹, we can determine the impact of *Mtb* coinfection on the course of malaria by daily monitoring parasitemia in Giemsa-stained blood smears as shown in **Figure 3B**.

Infection of mice with *Mtb* by the use of an inhalation exposure system results in successful deposition of bacteria in the lungs with relatively low variability between individual mice (**Figure 4A**). We can follow mycobacterial loads in lungs and other organs of interest over time by CFU determination. An example for the mycobacterial load in lungs of *Mtb* infected C57BL/6 mice in the absence or presence of *P. berghei* is shown in **Figure 4B**.

Table 1. Prepatency after sporozoite transmission by mosquito bite. All mice developed blood-stage infections after natural transmission of *P. berghei* sporozoites by mosquito bite. Adapted from¹ *PLoS ONE*, 7(10), e48110 with permission.

Experimental mouse group (C57BL/6)	Challenge by 10 infectious mosquito bites	No. of blood-stage positive animals / No. of animals per group	Mean prepatency [d]
naïve	<i>P. berghei</i>	7/7	4,4
<i>Mtb</i> infected	<i>P. berghei</i>	7/7	5,5

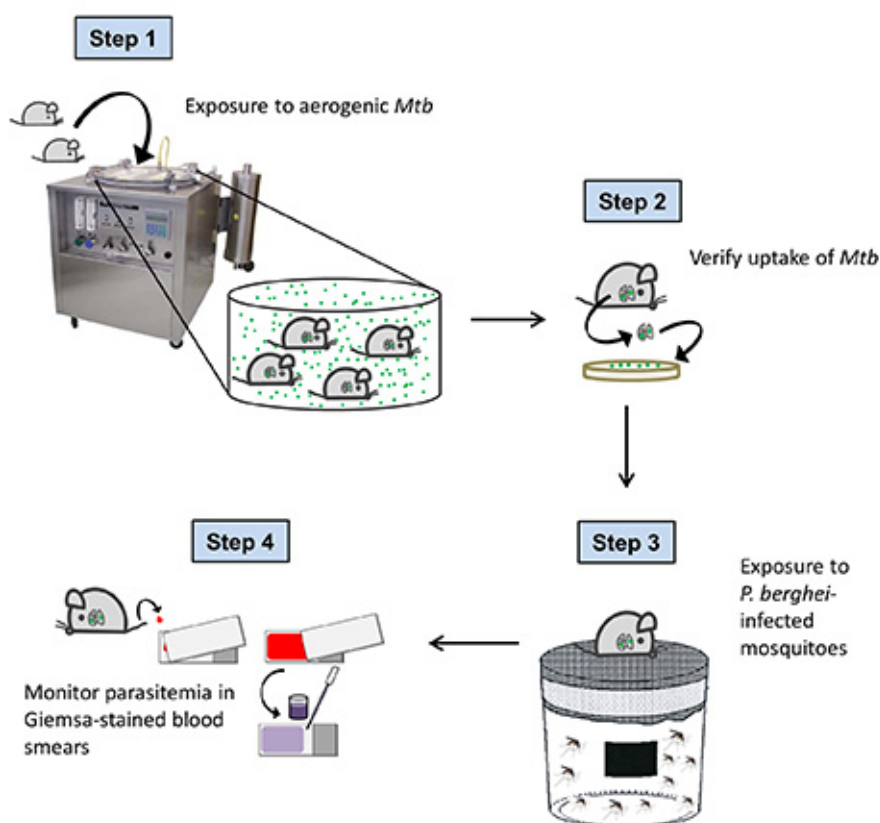


Figure 1. Overall scheme of the experimental procedure. C57BL/6 mice are exposed to *Mtb*-containing aerosol droplets in a Glas-Col aerosol chamber. One day after aerosol infection, the uptake of *Mtb* is verified by CFU analysis of the lung. 40 days after *Mtb* challenge (when tuberculosis has become chronic in infected mice), animals are exposed to *P. berghei*-infected mosquitoes. To confirm successful sporozoite transmission and to follow the course of *Plasmodium* infection, parasitemia is monitored daily in Giemsa-stained blood smears starting 3 days after mosquito bite.

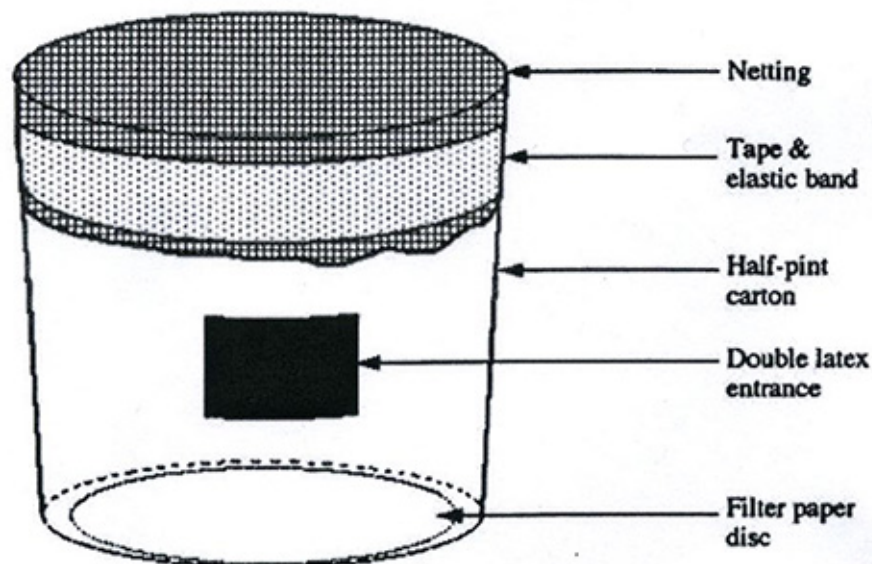


Figure 2. Sketch of a mosquito cage. Infectious mosquitoes are contained inside a carton such as a half-pint ice cream carton. A small opening is cut into the side of the carton and covered with a double layer of latex with a slit cut in each piece to create a secure entrance for the mosquitoes. A piece of filter paper is taped on the inside bottom of the carton to soak up drips. Ice cream cups are closed off with netting (Nylon mesh) which is fixed to the carton by tape and elastic band.

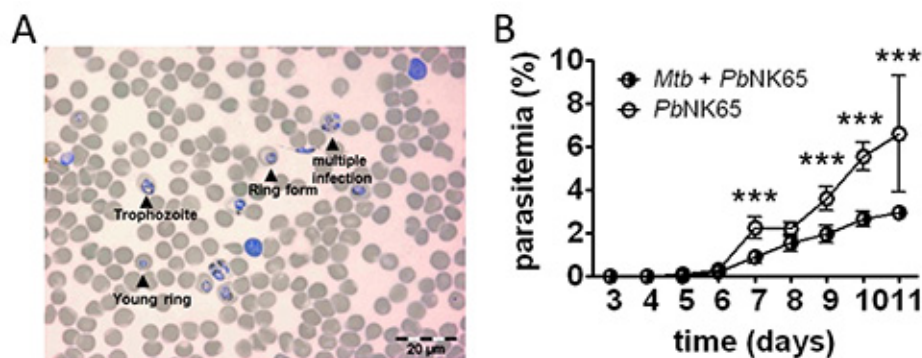


Figure 3. Blood parasitemia. **A)** Example of a Wright's stained blood smear showing parasitized erythrocytes (Scale bar, 20 μ m). **B)** Parasitemia in mouse blood was monitored by analyzing daily Giemsa-stained blood smears. Coinfected mice have a reduced parasitemia compared to *Plasmodium* single infected mice (n=10); results are shown as means \pm SD; ***p<0,001. Reprinted from ¹ PLoS ONE, 7(10), e48110 with permission.

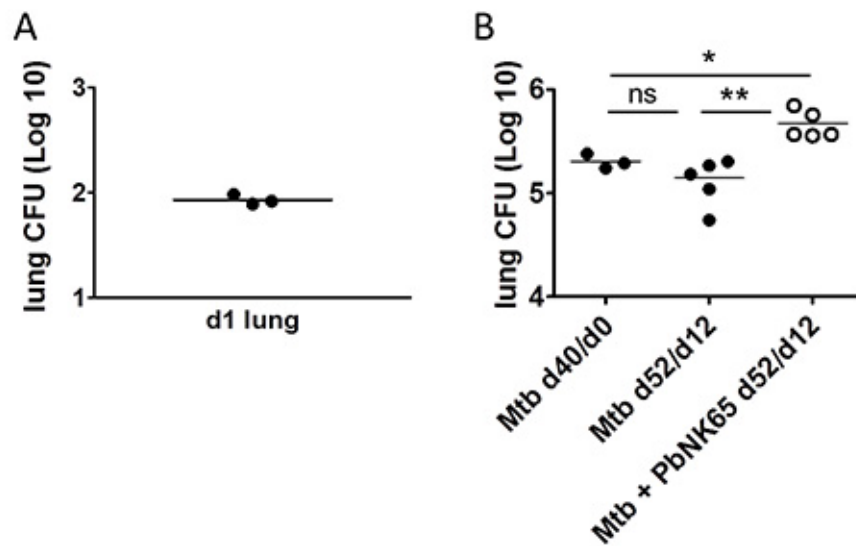


Figure 4. Detection of *Mtb* in the lung. C57BL/6 mice were aerosol challenged with 100 CFU of *Mtb* and 40 days later, coinfecting with *P. berghei* by mosquito bite. **A)** One day after aerosol infection the uptake of the desired number of *Mtb* was verified by plating whole lung homogenates for CFU determination. **B)** Bacterial loads at the time of coinfection were determined in lung lysates of 3 mice via CFU analysis (*Mtb* d40/d0). 12 days after coinfection, lung lysates were plated for CFU analysis to determine the impact of *Plasmodium* coinfection on *Mtb* control. Note, that *Mtb* numbers in lungs increased during coinfection with *P. berghei* NK65. X-axis labeling: time after *Mtb*-infection/time after coinfection. Adapted from¹ *PLoS ONE*, 7(10), e48110 with permission.

Discussion

We described how mice can be productively infected with *Mtb* and *P. berghei* via their natural routes of infection. We apply established infection protocols to study experimental tuberculosis¹³ or malaria¹² and recently adopted them to study coinfection between *Mtb* and *Plasmodium* in the mouse model¹.

The most critical steps for successful infections are the transmission of both pathogens at the desired numbers. Mycobacterial stocks should not be older than 2 years because they will lose viability over time, and the originally determined titer of the stocks will most likely no longer be accurate. As a result, infection doses would be much lower than expected. Therefore, CFU titers of stock cultures should be determined every few months. Equally important is the standardized cultivation and generation of *Mtb* stocks in the first place. Culture conditions such as medium, supplements, time, and volume should be standardized in order to be able to compare data from experiments using different stocks of *Mtb*. Mycobacteria tend to clump during culture, therefore it is important to keep culture conditions standardized, harvest bacteria at the same growth phase and resuspend frequently during aliquoting. Otherwise there can be a wide variation in CFU titers and infection outcome.

The generation of homogeneously infected mosquito batches is another critical step. Since different mosquitoes will feed on each individual mouse, it is important that all of the mosquitoes used for one infection experiment come from the same batch and that only those batches where more than 90% of mosquitoes are infected are used.

The current model can be used to dissect immune responses and immune modulations in a coinfecting host by comparing it to the immune responses in single infected hosts. We can apply various well established methodologies such as histology, immunohistochemistry, cytometry analysis of immune cell populations, or PCR and ELISA for cytokine and chemokine detection to the tissue or body fluids of interest. It should be noted, that such mouse models have certain limitations. Whereas the majority of people infected with *Mtb* develop a latent infection with no signs of clinical symptoms, mice develop a chronic disease with progressive lung pathology. Still, C57BL/6 mice are relatively resistant to *Mtb* infection and do not develop classical granulomas as observed in human tuberculosis patients. As a result, immunopathological reactions in the mouse do reflect neither latent nor active human tuberculosis disease appropriately. Despite the discrepancy between latent and chronic disease in man and mouse, mice have been extensively used to identify and study host factors that control *Mtb* infection and are valuable and indispensable tools to investigate immune responses in infectious diseases. Importantly, susceptibility to *Mtb* infection varies considerably between commonly used inbred mouse strains and more susceptible strains such as DBA or C3H can be included in co- and single-infection studies. Moreover, apart from the pathogen species described herein, any other *Mycobacterium* species of interest (e.g. clinical isolates) could be used. Likewise, while no single malaria model replicates all aspects of human disease different mouse models do closely resemble different aspects of naturally acquired human malaria infection. For instance, *P. berghei* ANKA and *P. yoelii* YM/17XL are widely studied models of human (*P. falciparum*-induced) severe malaria, with *P. berghei* ANKA being regarded as an excellent model for human cerebral malaria^{14,15}. *P. berghei* NK65 is an excellent model for hyperparasitaemia and acute malarial anemia, and has recently been described as an experimental model for malaria-associated acute respiratory distress syndrome (MA-ARDS¹⁶). Using the described coinfection model, we could recently show that concurrent *P. berghei* NK65 infection exacerbates chronic tuberculosis¹.

Apart from using different rodent malaria parasites, the order and timing of infection events may be adapted depending on the underlying research question. In the field, concomitant *Plasmodium* infections may be present at the time of *Mtb* infection or acquired subsequently. In both scenarios the immune responses to *Mtb* and *Plasmodium* may be affected differently. Therefore, mice can be infected with *Plasmodium* sporozoites either before or after *Mtb* infection. Furthermore, mice can be challenged at different times post *Mtb* infection to assess the impact of the *Plasmodium*-induced immune response on acute versus chronic tuberculosis. Importantly, when choosing a malaria parasite for studying malaria-tuberculosis coinfection, one should be aware that some rodent *Plasmodium* strains cause acute disease in certain mouse strains and succumb to infection within only days or weeks while *Mtb* causes chronic and long-lasting infections in immunocompetent mice. Hence, timing of coinfection is crucial in order to avoid premature death of experimental animals.

The modulation of host immunity by simultaneous infection with multiple pathogen species remains poorly understood. Our experimental coinfection model provides a powerful tool to investigate *Mycobacterium-Plasmodium* interaction with the immune system of a coinfecting host and will contribute to our understanding how coinfections can affect pathogenesis, disease outcome, vaccination, immunodiagnostics, and therapy.

Disclosures

The authors have nothing to disclose.

Acknowledgements

The authors would like to thank Miriam Ester for mosquito breeding. This work was supported by in-house funding from the Research Center Borstel and joined funding by the Leibniz Center Infection.

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